CLAIMS

- A method of producing endostatin protein comprising:
 recombinantly producing endostatin using an expression system;
 recovering endostatin; and,
 purifying endostatin.
- 2. The method of Claim 1, further comprising lyophilizing endostatin.
- 3. The method of Claim 1, wherein the expression system is *Pichia pastoris*, yeast, *E. coli*, insect cells, baculovirus, transgenic animals, or transgenic plants.
- 4. The method of Claim 1, wherein the expression system is *Pichia pastoris*.
- 5. The method of Claim 1, wherein the recombinantly produced endostatin has an amino acid sequence shown in SEQ ID NO: 3, SEQ ID NO: 5, SEQ ID NO: 6, SEQ ID NO: 7, SEQ ID NO: 8, or SEQ ID NO: 11, or a fragment thereof.
- 6. The method of Claim 1, wherein the recombinantly produced endostatin is encoded by a nucleotide sequence shown in SEQ ID NO: 4 or SEQ ID NO: 10, or a fragment thereof.
- 7. A method of recombinantly producing endostatin protein comprising: preparing an inoculum culture; and fermenting the culture.
- 8. The method of Claim 7, wherein the inoculum culture is a two stage seed process of *Pichia pastoris*.

- 9. The method of Claim 7, wherein the fermenting step includes fermentation media comprising calcium sulfate, potassium sulfate, magnesium sulfate, potassium hydroxide, phosphoric acid and glycerol.
- 10. The method of Claim 7, wherein the fermenting step comprises a fermentation process comprising a batch glycerol phase, a fed-batch glycerol phase, a methanol ramp phase and methanol induction phase.
- 11. A method of purifying endostatin protein comprising:

capturing endostatin from a sample using a first cation exchange column and expanded bed chromatography;

applying the endostatin to a heparin-sepharose column or to a column containing a resin useful for hydrophobic interaction chromatography;

applying the endostatin to an anion exchange column; applying the endostatin to a second cation exchange column; and, concentrating the endostatin.

- 12. The method of Claim 11, wherein the resin useful for hydrophobic interaction chromatography is phenyl sepharose resin.
- 13. The method of Claim 11, wherein the anion exchange column is an amine column.
- 14. The method of Claim 11, wherein first cation exchange column contains STREAMLINETM sulfopropyl resin or carboxymethylcellulose.
- 15. The method of Claim 11, wherein concentrating the endostatin further comprises pushing the sample through a membrane containing a molecular weight cutoff selected for endostatin and eluting endostatin from the membrane with buffer.
- 16. The method of Claim 15, wherein the eluted endostatin is lyophilized.

- 17. The method of Claim 15, wherein the membrane is made from polyethersulfone.
- 18. The method of Claim 11, wherein concentrating the endostatin further comprises use of parallel flow concentrators.
- 19. The method of Claim 15, wherein the buffer comprises a citrate-phosphate buffer.
- 20. The method of Claim 19, further comprising removal of citrate by exchanging with phosphate buffered saline and a detergent.
- 21. The method of Claim 20, further comprising lyophilizing endostatin.
- 22. The method of Claim 21, further comprising reconstituting the lyophilized endostatin with a solution.
- 23. The method of Claim 22, wherein the solution is an aqueous zinc chloride solution.